L Number	Hits	Search Text	DB	Time stamp
1	611	(alzheimer same disease) and (cholinesterase same inhibitor)	USPAT;	2004/04/22 11:21
ŀ			US-PGPUB;	
			EPO; JPO;	
			DERWENT	
3	0	(alzheimer same disease) same (cholinesterase same inhibitor) same treat?	USPAT;	2004/04/22 11:22
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		·	EPO; JPO;	
			DERWENT	
-	2	daly-james.in.	USPAT;	2004/04/22 11:21
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			EPO; JPO;	
			DERWENT	
-	2	kotwal-girish.in.	USPAT;	2004/04/22 10:22
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			EPO; JPO;	
		,	DERWENT	
-	46	vaccinia same complement same control same protein	USPAT;	2004/04/22 10:24
İ			US-PGPUB;	
			EPO; JPO;	
			DERWENT	
-	4	(vaccinia same complement same control same protein) and alzheimer	USPAT;	2004/04/22 10:27
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			EPO; JPO;	
			DERWENT	
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			EPO; JPO;	
			DERWENT	

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MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
AND CURRENT DISCOVER FILE IS DATED 13 APRIL 2004

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FILE 'HOME' ENTERED AT 10:35:39 ON 22 APR 2004

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=> s daly james/au

L117 DALY JAMES/AU

=> s kotwal girish/au

0 KOTWAL GIRISH/AU L₂

=> s (vaccinia (s) complement (s) control (s) protein) (p) alzheimer 15 (VACCINIA (S) COMPLEMENT (S) CONTROL (S) PROTEIN) (P) ALZHEIMER

=> dup rem 13

PROCESSING COMPLETED FOR L3

6 DUP REM L3 (9 DUPLICATES REMOVED)

=> d 14 total ibib kwic

ANSWER 1 OF 6 DUPLICATE 1 MEDLINE on STN

ACCESSION NUMBER: 2004003750 IN-PROCESS

PubMed ID: 14698003 DOCUMENT NUMBER:

Prolonged retention of vaccinia virus complement control TITLE: protein following IP injection: implications in blocking

xenorejection.

AUTHOR: Jha P; Smith S A; Justus D E; Kotwal G J

Department of Microbiology and Immunology, University of CORPORATE SOURCE:

Louisville School of Medicine, Louisville, KY 40202, USA.

Transplantation proceedings, (2003 Dec) 35 (8) 3160-2. SOURCE:

Journal code: 0243532. ISSN: 0041-1345.

PUB. COUNTRY: United States

Journal; Article; (JOURNAL ARTICLE) DOCUMENT TYPE:

LANGUAGE: English

FILE SEGMENT: IN-PROCESS; NONINDEXED; Priority Journals

ENTRY DATE: Entered STN: 20040106

Last Updated on STN: 20040129

The vaccinia virus complement control AB

> protein (VCP) blocks classic and alternate complement pathways by binding to the third and fourth complement components and by blocking the formation of the C3-convertase as well as by accelerating the decay of the C3 and C4 convertase. The therapeutic potential of VCP has been extensively studied for brain injury, xenotransplantation, Alzheimer's disease, and spinal cord injury. We investigated the pharmacokinetic behavior of rVCP in mice. Dosage of rVCP was studied by.

ANSWER 2 OF 6 MEDLINE on STN DUPLICATE 2 L4

ACCESSION NUMBER: 2003213362 MEDLINE DOCUMENT NUMBER: PubMed ID: 12734405

TITLE: Vaccinia complement control protein: multi-functional

protein and a potential wonder drug.

AUTHOR: Jha Purushottam; Kotwal Girish J

CORPORATE SOURCE: Department of Microbiology and Immunology, University of

Louisville, School of Medicine, Louisville, KY 40202, USA. Journal of biosciences, (2003 Apr) 28 (3) 265-71. Ref: 36 Journal code: 8100809. ISSN: 0250-5991.

PUB. COUNTRY:

SOURCE:

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

General Review; (REVIEW)

(REVIEW, TUTORIAL)

LANGUAGE:

English

FILE SEGMENT:

Priority Journals

ENTRY MONTH:

200403

ENTRY DATE:

Entered STN: 20030508

Last Updated on STN: 20040324 Entered Medline: 20040323

AB Vaccinia virus complement control

protein (VCP) was one of the first viral molecules demonstrated to
have a role in blocking complement and hence in the evasion of
host defense. Structurally it is very similar to the human C4b-BP and the
other. . . can take place simultaneously and contribute to its many
function and to its potential use in several inflammatory diseases, e.g.
Alzheimer's disease (AD), CNS injury, xenotransplantation, etc.
making it a truly fascinating molecule and potential drug.

L4 ANSWER 3 OF 6 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN

ACCESSION NUMBER:
DOCUMENT NUMBER:

2003:145171 BIOSIS PREV200300145171

TITLE:

Potential intervention by vaccinia virus

complement control protein of

the signals contributing to the progression of central

nervous system injury to Alzheimer's disease.

AUTHOR (S):

Kotwal, Girish J. [Reprint Author]; Lahiri, Debomoy K.;

Hicks, Ramona

CORPORATE SOURCE:

Department of Microbiology and Immunology, University of Louisville School of Medicine, Louisville, KY, 40202, USA

gjkotw01@gwise.louisville.edu

SOURCE:

Diederich, Marc [Editor, Reprint Author]. (2002) pp.

317-322. Cell signaling, transcription, and translation as

therapeutic targets. print.

Publisher: New York Academy of Sciences, 2 East 63rd

Street, New York, NY, 10021, USA. Series: Annals of the New

York Academy of Sciences.

Meeting Info.: Conference on Cell Signaling, Transcription

and Translation as Therapeutic Targets. Luxembourg,

Luxembourg. January 30-February 02, 2002. University Center

Luxembourg, Department of Sciences; Doctoral School Medecine et Sante; University Henri Poincare Nancy I;

National Research Fund.

ISSN: 0077-8923 (ISSN print). ISBN: 1-57331-428-5 (cloth),

1-57331-429-3 (paper).

DOCUMENT TYPE:

Book; (Book Chapter) Conference; (Meeting)

Conference; (Meeting Paper)

LANGUAGE:

English

ENTRY DATE:

Entered STN: 19 Mar 2003

Last Updated on STN: 19 Mar 2003

TI Potential intervention by vaccinia virus complement control protein of the signals contributing to the

progression of central nervous system injury to Alzheimer's

disease.

L4 ANSWER 4 OF 6

MEDLINE on STN

DUPLICATE 3

ACCESSION NUMBER:

2002729264 MEDLINE

DOCUMENT NUMBER: PubMed ID: 12485887

TITLE:

Potential intervention by vaccinia virus

complement control protein of

the signals contributing to the progression of central

nervous system injury to Alzheimer's disease.

AUTHOR: CORPORATE SOURCE: Kotwal Girish J; Lahiri Debomoy K; Hicks Ramona Department of Microbiology and Immunology, University of

Louisville School of Medicine, Louisville, Kentucky 40202,

USA.. gjk01@gwise.louisville.edu

SOURCE:

Annals of the New York Academy of Sciences, (2002 Nov) 973

317-22. Ref: 40

Journal code: 7506858. ISSN: 0077-8923.

PUB. COUNTRY:

United States

DOCUMENT TYPE:

Journal; Article; (JOURNAL ARTICLE)

General Review; (REVIEW)

(REVIEW, TUTORIAL)

LANGUAGE:

English

FILE SEGMENT:

Priority Journals

ENTRY MONTH:

200302

ENTRY DATE:

Entered STN: 20021221

Last Updated on STN: 20030214 Entered Medline: 20030213

TI Potential intervention by vaccinia virus complement control protein of the signals contributing to the progression of central nervous system injury to Alzheimer's disease.

Alzheimers disease (AD) and for depression. The mechanisms by which trauma causes delayed cognitive deficits are largely unknown. In recent studies,. . . and other previous studies, it was hypothesized that regulation of the complement system will attenuate the long-term consequences of TBI. Vaccinia virus complement control protein (VCP) is a protein encoded by vaccinia virus. It blocks both the classic and alternative pathways of complement activation in vitro, and by doings so prevents the.

L4 ANSWER 5 OF 6 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1999:212578 CAPLUS

DOCUMENT NUMBER:

131:57660

TITLE:

Pro-inflammatory complement activation by the $A\beta$ peptide of Alzheimer's disease

is biologically significant and can be blocked by

vaccinia virus complement

control protein

AUTHOR(S):

Daly, James; Kotwal, Girish J.

CORPORATE SOURCE:

Department of Microbiology and Immunology, University

of Louisville School of Medicine, Louisville, KY,

40292, USA

SOURCE:

Neurobiology of Aging (1999), Volume Date 1998, 19(6),

619-627

CODEN: NEAGDO; ISSN: 0197-4580

PUBLISHER:

Elsevier Science Inc.

DOCUMENT TYPE: LANGUAGE:

Journal English

REFERENCE COUNT:

THERE ARE 35 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

TI Pro-inflammatory complement activation by the Aβ peptide of
Alzheimer's disease is biologically significant and can be blocked
by vaccinia virus complement control
protein

The amyloid plaque is the hallmark of Alzheimer's disease (AD). The transmembrane domain and a portion of the C-terminus (A β) of the amyloid precursor protein, are known to form the nucleus of the amyloid plaque. It has been demonstrated recently, using in vitro assays, that the A β peptide can activate both the classical (antibody-independent) and alternate pathways of complement activation. The proposed complement activation is due to the binding of A β to the complement components Clq and C3, resp., which initiate formation of the proinflammatory C5a and C5b-9 membrane attack complex. In this report, the authors have investigated the in vitro findings for the likely complement-dependent proinflammatory properties of the Alzheimer's disease A β peptide. The authors have performed expts. using congenic C5-deficient and C5-sufficient mice injected with synthetic A β and recombinant

```
polypeptide (C-100) containing Aβ. Injection of C-100 into C5-sufficient
mice induced a clear increase in the number of polymorphonuclear cells
(neutrophils) at the site of injection due to complement activation and
the subsequent release of proinflammatory chemtoactic factors. In sharp
contrast, the C5-deficient mice did not show any increase in cellular
influx. The vaccinia virus complement control
protein, an inhibitor of both the classical and alternate pathway
can down-regulate the biol. significant activation of complement
by A\beta, as demonstrated by an in vitro immunoassay. The therapeutic
down-regulation of Aβ-caused complement activation could greatly
alleviate the progression of some of the chronic neurodegeneration
characteristic of Alzheimer's disease.
Proteins, specific or class
RL: BAC (Biological activity or effector, except adverse); BSU (Biological
study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
   (VCP (vaccinia virus complement control
  protein); pro-inflammatory complement activation by
   Aß peptide of Alzheimer's disease is biol. significant
   and can be blocked by vaccinia virus complement
   control protein)
Complement
   (activation; pro-inflammatory complement activation by
   Aß peptide of Alzheimer's disease is biol. significant
   and can be blocked by vaccinia virus complement
   control protein)
Complement
RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)
   (alternative pathway; pro-inflammatory complement activation
   by A\beta peptide of Alzheimer's disease is biol.
   significant and can be blocked by vaccinia virus
   complement control protein)
Protein sequences
cDNA sequences
   (amyloid precursor protein C-terminal fragment;
   pro-inflammatory complement activation by A\beta peptide of
   Alzheimer's disease is biol. significant and can be blocked by
   vaccinia virus complement control
   protein)
Alzheimer's disease
Inflammation
  Vaccinia virus
   (pro-inflammatory complement activation by Aβ peptide of
   Alzheimer's disease is biol. significant and can be blocked by
   vaccinia virus complement control
   protein)
Amyloid precursor proteins
RL: ADV (Adverse effect, including toxicity); PRP (Properties); BIOL
(Biological study)
   (pro-inflammatory complement activation by Aβ peptide of
   Alzheimer's disease is biol. significant and can be blocked by
   vaccinia virus complement control
   protein)
Brain, disease
   (senile plaque; pro-inflammatory complement activation by
   Aβ peptide of Alzheimer's disease is biol. significant
   and can be blocked by vaccinia virus complement
   control protein)
Amyloid
RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)
   (\beta-; pro-inflammatory complement activation by A\beta
   peptide of Alzheimer's disease is biol. significant and can
   be blocked by vaccinia virus complement
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control protein)

134548-35-9, 652-751-Glycoprotein (human clone λΑΡCP168i4 amyloid ITA4 precursor protein moiety reduced) RL: PRP (Properties) (amino acid sequence; pro-inflammatory complement activation by Aβ peptide of Alzheimer's disease is biol. significant and can be blocked by vaccinia virus complement control protein) 228092-51-1 ITRL: PRP (Properties) (nucleotide sequence; pro-inflammatory complement activation by $A\beta$ peptide of Alzheimer's disease is biol. significant and can be blocked by vaccinia virus complement control protein) 80295-53-0, Complement C5 IT RL: ADV (Adverse effect, including toxicity); BIOL (Biological study) (pro-inflammatory complement activation by $A\beta$ peptide of Alzheimer's disease is biol. significant and can be blocked by vaccinia virus complement control protein) DUPLICATE 4 ANSWER 6 OF 6 MEDLINE on STN ACCESSION NUMBER: MEDLINE 1999206442 PubMed ID: 10192224 DOCUMENT NUMBER: Pro-inflammatory complement activation by the A TITLE: beta peptide of Alzheimer's disease is biologically significant and can be blocked by vaccinia virus complement control protein. Daly J 4th; Kotwal G J AUTHOR: Department of Microbiology and Immunology, University of CORPORATE SOURCE: Louisville School of Medicine, KY 40292, USA. Neurobiology of aging, (1998 Nov-Dec) 19 (6) 619-27. SOURCE: Journal code: 8100437. ISSN: 0197-4580. PUB. COUNTRY: United States Journal; Article; (JOURNAL ARTICLE) DOCUMENT TYPE: LANGUAGE: English Priority Journals FILE SEGMENT: ENTRY MONTH: 199905 ENTRY DATE: Entered STN: 19990525 Last Updated on STN: 19990525 Entered Medline: 19990511 Pro-inflammatory complement activation by the A beta peptide of TTAlzheimer's disease is biologically significant and can be blocked by vaccinia virus complement control protein. AΒ The amyloid plaque is the hallmark of Alzheimer's disease (AD). The transmembrane domain and a portion of the C-terminus (A beta) of the amyloid precursor protein, are known. . . attack complex. In this report, we have investigated the in vitro findings for the likely complement-dependent proinflammatory properties of the Alzheimer 's disease A beta peptide. We have performed experiments using congenic C5-deficient and C5-sufficient mice injected with synthetic A beta and. release of proinflammatory chemtoactic factors. In sharp contrast, the C5-deficient mice did not show any increase in cellular influx. The vaccinia virus complement control protein, an inhibitor of both the classical and alternate pathway can down-regulate the biologically significant activation of complement by A beta, as demonstrated by an in vitro immunassay. The therapeutic down-regulation of A beta-caused complement activation could greatly alleviate the progression of some of the chronic

neurodegeneration characteristic of Alzheimer's disease.

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09889624 Results

SEQ ID NO: 1

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(USDC) US SEC OF COMMERCE.

SUMMARIES

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PR
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XX
     (USSH ) NAT INST OF HEALTH.
PA
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ΡI
    Kotwal G;
хx
    WPI; 1989-165451/22.
DR
    N-PSDB; AAN90113.
DR
ХХ
    New protein with anti-complement activity - encoded by Vaccinia virus 35K
PT
PT
XX
    Disclosure; Fig 2A; 20pp; English.
PS
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    C4b-binding protein which specifically blocks human complement cascades.
CC
    It is the deduced sequence of a 35kDa protein encoded by sequence 52-840
CC
    of the 35K gene of vaccinia virus strain WR. Note a = these sites
CC
    indicate the start of 60 amino acid tandem repeating units which have a
CC
    consensus sequence. The signal peptide sequence is not found in purified
CC
    35K protein recovered from the medium of cells infected with vaccinia
CC
    virus strain WR. A suggested use is to treat diseases due to abnormally
    high complement activity. (Note: Revised entry submitted to correct the
CC
    patent number format of US Government-owned NTIS applications to prevent
CC
    clashes with ongoing US granted patent numbers. For further information
CC
    please visit the Derwent web site at
CC
    www.derwent.com/dwpi/updates/ntis_us.html.) (Updated on 10-MAR-2003 to
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XX
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5	706	48.2	126	6	5514582-43	Patent No. 5514582
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7	407	27.8	2489	4	US-09-911-842A-5	Sequence 5, Appli
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RESULT 1
US-09-653-813-2
; Sequence 2, Application US/09653813
; Patent No. 6551595
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 APPLICANT: ROSENGARD, Ariella M.
  APPLICANT: AHEARN, Joseph M.
  TITLE OF INVENTION: SMALLPOX INHIBITOR OF COMPLEMENT ENZYMES (SPICE)
  TITLE OF INVENTION: PROTEIN AND METHODS OF INHIBITING COMPLEMENT ACTIVATION
  FILE REFERENCE: 9596-107U1
  CURRENT APPLICATION NUMBER: US/09/653,813
  CURRENT FILING DATE: 2000-09-01
  PRIOR APPLICATION NUMBER: US 60/076,821
  PRIOR FILING DATE: 1998-03-03
  PRIOR APPLICATION NUMBER: WO PCT/US99/04635
  PRIOR FILING DATE: 1999-03-02
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US-09-653-813-2
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RESULT 2
US-07-906-983-2
; Sequence 2, Application US/07906983
 Patent No. 5187268
  GENERAL INFORMATION:
    APPLICANT: Kotwal, Girish
    APPLICANT: Moss, Bernard
    TITLE OF INVENTION: Synthetic, Anti-Complement Protein and
    TITLE OF INVENTION: the Gene Encoding Same
    NUMBER OF SEQUENCES: 3
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: Townsend and Townsend
      STREET: One Market Plaza, Steuart Tower, Suite 2000
      CITY: San Francisco
      STATE: California
      COUNTRY: USA
      ZIP: 94105
    COMPUTER READABLE FORM:
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MEDIUM TYPE: Floppy disk

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COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.25
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/07/906,983
      FILING DATE: 19920701
      CLASSIFICATION: 530
    ATTORNEY/AGENT INFORMATION:
      NAME: Weber, Kenneth A.
      REGISTRATION NUMBER: 31,677
      REFERENCE/DOCKET NUMBER: 15280-9
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: 415-543-9600
      TELEFAX: 415-543-5043
  INFORMATION FOR SEQ ID NO: 2:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 263 amino acids
      TYPE: AMINO ACID
      TOPOLOGY: linear
    MOLECULE TYPE: protein
US-07-906-983-2
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                                            2; Indels
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                        263 2 T28450
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                        263 2 B72152
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                        597 1 NBHUC4
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          467
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    9
                        558 2 S57953
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    10
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                29.9
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          432
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A: Note: host Homo sapiens (man)
C;Date: 31-Dec-1989 #sequence_revision 30-Jun-1990 #text_change 27-Oct-2003
C; Accession: A31005; B42504
R; Kotwal, G.J.; Moss, B.
Nature 335, 176-178, 1988
A; Title: Vaccinia virus encodes a secretory polypeptide structurally related to
complement control proteins.
A; Reference number: A31005; MUID: 88318974; PMID: 3412473
A; Accession: A31005
A; Molecule type: DNA
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A; Cross-references: GB:X13166; NID:g60690; PIDN:CAA31564.1; PID:g60691
A; Experimental source: strain WR
R; Goebel, S.J.; Johnson, G.P.; Perkus, M.E.; Davis, S.W.; Winslow, J.P.; Paoletti, E.
Virology 179, 517-563, 1990
A; Title: Appendix to "The complete DNA sequence of vaccinia virus".
A; Reference number: A42501
A:Accession: B42504
A; Molecule type: DNA
A; Residues: 1-263 <GOE>
A; Cross-references: GB: M35027; NID: g335317; PIDN: AAA47997.1; PID: g335345
A; Experimental source: strain Copenhagen
R; Goebel, S.J.; Johnson, G.P.; Perkus, M.E.; Davis, S.W.; Winslow, J.P.; Paoletti, E.
Virology 179, 247-266, 1990
A; Title: The complete DNA sequence of vaccinia virus.
A; Reference number: A42531; MUID: 91021027; PMID: 2219722
A; Contents: annotation; possible protein-coding frames
A; Note: neither amino acid nor nucleotide sequence is given
C; Superfamily: complement control protein; complement factor H repeat homology
C; Keywords: duplication; extracellular protein
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F;21-81/Domain: complement factor H repeat homology <FH1>
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F;148-201/Domain: complement factor H repeat homology <FH3>
F;206-261/Domain: complement factor H repeat homology <FH4>
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C; Species: variola major virus
C;Date: 22-Oct-1999 #sequence_revision 22-Oct-1999 #text_change 27-Oct-2003
C:Accession: T28450
R; Massung, R.F.; Esposito, J.J.; Liu, L.I.; Qi, J.; Utterback, T.R.; Knight, J.C.; Aubin,
L.; Yuran, T.E.; Parsons, J.M.; Loparev, V.N.
Nature 366, 748-751, 1993
A; Title: Potential virulence determinants in terminal regions of variola smallpox virus
genome.
A; Reference number: Z20488; MUID: 94088747; PMID: 8264798
A; Accession: T28450
A;Status: preliminary; translated from GB/EMBL/DDBJ
A; Molecule type: DNA
A; Residues: 1-263 < MAS>
A;Cross-references: EMBL:L22579; NID:g623595; PIDN:AAA60760.1; PID:g438930
A; Experimental source: strain "Bangladesh-1975"
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Qу
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                 26.7
                        1033 1 CR2 HUMAN
                                                           P20023 homo sapien
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     01-JUL-1989 (Rel. 11, Created)
     01-JUL-1989 (Rel. 11, Last sequence update)
     10-OCT-2003 (Rel. 42, Last annotation update) ·
     Complement control protein precursor (VCP) (Secretory protein 35)
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     Vaccinia virus (strain WR), and
     Vaccinia virus (strain Copenhagen).
     Viruses; dsDNA viruses, no RNA stage; Poxviridae; Chordopoxvirinae;
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     STRAIN=WR;
     MEDLINE=88318974; PubMed=3412473;
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     to complement control proteins.";
     Nature 335:176-178(1988).
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     STRAIN=WR;
    MEDLINE=89073756; PubMed=2849238;
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     Virology 167:524-537(1988).
     [3]
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    MEDLINE=91021027; PubMed=2219722;
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     Virology 179:247-266(1990).
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    STRAIN=Copenhagen;
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    MEDLINE=92115714; PubMed=1731333;
    Isaacs S.N., Kotwal G.J., Moss B.;
    "Vaccinia virus complement-control protein prevents
    antibody-dependent complement-enhanced neutralization of infectivity
    and contributes to virulence.";
    Proc. Natl. Acad. Sci. U.S.A. 89:628-632(1992).
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ID

AC

DТ

DT

DE DE

GN

OS

os

OC

OC

OX

RN

RP

RC

RX RA

RT

RT

RL

RN

RP

RC

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RL

RN

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RA RT

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RΑ
     Barlow P.N.;
     "NMR studies of a viral protein that mimics the regulators of
RT
     complement activation.";
RL
     J. Mol. Biol. 272:253-265(1997).
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CC
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CC
     This SWISS-PROT entry is copyright. It is produced through a collaboration
CC
     between the Swiss Institute of Bioinformatics and the EMBL outstation -
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     the European Bioinformatics Institute. There are no restrictions on its
     use by non-profit institutions as long as its content is in no way
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     or send an email to license@isb-sib.ch).
CC
DR
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FΤ
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FT
                155
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                157
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FT
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                       176
FT
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6	1385	94.5	263	12	Q89076	Q89076 variola vir
7	1385	94.5	263	12	Q07033	Q07033 variola vir
8	1384	94.4	259	12	P87616	P87616 cowpox viru
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13	491.5	33.5	365	6	Q7YRJ3	Q7yrj3 bos taurus
14	486.5	33.2	679	11	Q99254	Q99254 mus musculu
15	466.5	31.8	645	12	Q9WRU2	Q9wru2 macaca mula
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RESULT 7
Q07033
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TD
                                    PRT:
                                           263 AA.
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DT
DT
     01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
     01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DT
DE
     D12L protein.
GN
     D12L.
os
     Variola virus.
OC
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OC
     Orthopoxvirus.
OX
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RP
     SEQUENCE FROM N.A.
     STRAIN=India-1967;
RC
RA
     Blinov V.M.;
RL
     Submitted (NOV-1992) to the EMBL/GenBank/DDBJ databases.
RN
RP
     SEQUENCE FROM N.A.
     STRAIN=India-1967;
RC
RX
     MEDLINE=93202281; PubMed=8384129;
     Shchelkunov S.N., Blinov V.M., Sandakhchiev L.S.;
RA
     "Genes of variola and vaccinia viruses necessary to overcome the host
RT
RT
     protective mechanisms.";
     FEBS Lett. 319:80-83(1993).
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[3]
RP
    SEQUENCE FROM N.A.
    STRAIN=India-1967;
RC
RX
    MEDLINE=95159666; PubMed=7856312;
    Shchelkunov S.N., Blinov V.M., Resenchuk S.M., Totmenin A.V.,
RA
    Olenina L.V., Chirikova G.B., Sandakhchiev L.S.;
RA
    "Analysis of the nucleotide sequence of 53 kbp from the right terminus
RT
    of the genome of variola major virus strain India-1967.";
RT
    Virus Res. 34:207-236(1994).
RL
RN
    [4]
    SEQUENCE FROM N.A.
RP
    STRAIN=India-1967;
    MEDLINE=95320969; PubMed=7597802;
Shchelkunov S.N., Totmenin A.V.;
RX
RA
    "Two types of deletions in orthopoxvirus genomes.";
    Virus Genes 9:231-245(1995).
RL
RN
RP
    SEQUENCE FROM N.A.
    STRAIN=India-1967;
RC
    MEDLINE=96290243; PubMed=8725113;
    Shchelkunov S.N., Totmenin A.V., Sandakhchiev L.S.;
RA
    "Analysis of the nucleotide sequence of 23.8 kbp from the left
RT
RT
    terminus of the genome of variola major virus strain India-1967.";
    Virus Res. 40:169-183(1996).
RL
    EMBL; X69198; CAA48953.1; -.
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    PIR; C36838; C36838.
    HSSP; P10998; 1VVD.
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    Pfam; PF00084; sushi; 4.
DR
DR
    SMART; SM00032; CCP; 4.
SQ
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  Query Match
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                                                                      0:
 Matches 248; Conservative
                                                 Indels
                                                           0; Gaps
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Qy
                 Db
           1 MKVERVTFLTLLGIGCVLSCCTIPSRPINMTFKNSVETDANANYNIGDTIEYLCLPGYRK 60
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Qy
             61 QKMGPIYAKCTGTGWTLFNQCIKRRCPSPRDIDNGHLDIGGVDFGSSITYSCNSGYYLIG 120
Db
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Qу
              121 EYKSYCKLGSTGSMVWNPKAPICESVKCOLPPSISNGRHNGYNDFYTDGSVVTYSCNSGY 180
Db
         181 SLIGNSGVLCSGGEWSDPPTCQIVKCPHPTISNGYLSSGFKRSYSYNDNVDFKCKYGYKL 240
Qy
             DЬ
         181 SLIGNSGVLCSGGEWSNPPTCQIVKCPHPTILNGYLSSGFKRSYSYNDNVDFTCKYGYKL 240
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Qу
             Dh
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RN